

1999 Annual Report

# NARMS

National Antimicrobial Resistance Monitoring System: Enteric Bacteria



CENTERS FOR DISEASE CONTROL  
AND PREVENTION

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# National Antimicrobial Resistance Monitoring System For Enteric Bacteria

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### Summary

In 1999, 1499 non-Typhi *Salmonella* isolates, 166 *Salmonella* Typhi isolates, 375 *Shigella*, 292 *E. coli* O157 isolates, and 319 *Campylobacter* isolates from humans were tested by the National Antimicrobial Resistance Monitoring System (NARMS) for Enteric Bacteria. Twenty-six percent of non-Typhi *Salmonella* isolates were resistant to one or more antimicrobial agents. Among *Salmonella* serotype Typhimurium isolates, 49% were resistant to one or more antimicrobial agents. Twenty-eight percent of *Salmonella* Typhimurium isolates had the multidrug-resistant pattern characteristic of DT104; resistant to ampicillin, chloramphenicol, streptomycin, sulfamethoxazole, and tetracycline (ACSSuT). One *Salmonella* isolate was resistant to ciprofloxacin (*S. Senftenberg*). The percentage of non-Typhi *Salmonella* isolates with ciprofloxacin minimum inhibitory concentrations (MICs)  $\geq 0.25$   $\mu\text{g/ml}$  increased from 0.4% in 1996 to 1.0% in 1999. The percentage of non-Typhi *Salmonella* isolates with a ceftriaxone MIC  $\geq 16$   $\mu\text{g/ml}$  increased from 0.1% in 1996 to 2% in 1999. Among *S. Typhi* isolates, 29% were resistant to one or more antimicrobial agents. Among *Shigella* isolates, 91% were resistant to one or more antimicrobial agents. Among *E. coli* O157 isolates, 10% were resistant to one or more antimicrobial agents. Among all *Campylobacter* isolates, 53% were resistant to one or more antimicrobial agents. Among *Campylobacter jejuni* isolates, 54% were resistant to one or more antimicrobial agents; 18% were resistant to ciprofloxacin.



## Methods

NARMS was launched in 1996, within the framework of CDC's Emerging Infections Program's Epidemiology and Laboratory Capacity Program and the Foodborne Disease Active Surveillance Network (FoodNet) as a collaboration among CDC, the U.S. Food and Drug Administration (FDA)-Center for Veterinary Medicine, U. S. Department of Agriculture (USDA)-Food Safety and Inspection Service and Agricultural Research Service, and state and local health departments to monitor prospectively the antimicrobial resistance of human non-Typhi *Salmonella* and *Escherichia coli* O157 isolates. Testing of *Campylobacter* isolates was added in 1997, and testing of *Salmonella* Typhi and *Shigella* isolates was added in 1999. In 1999, there were 17 NARMS health department participants (CA, CO, CT, FL, GA, KS, Los Angeles County, MD, MN, MA, NJ, New York City, NY, OR, TN, WA, and WV), representing approximately 103 million persons (38% of the United States population), and 7 of the 9 U.S. regions [Table 1]. In 1999, seven states (CA, CT, GA, MD, MN, NY, and OR) also monitored antimicrobial resistance among human *Campylobacter* isolates.

In 1999, NARMS participating public health laboratories have selected every tenth non-Typhi *Salmonella*, every *Salmonella* Typhi, every tenth *Shigella*, and every fifth *E. coli* O157 isolate received at their laboratory, and forwarded the isolates to CDC for susceptibility testing. Although we requested that participating laboratories send every *S. Typhi* isolate, analysis was restricted to one isolate per patient. At CDC, a semiautomated system (Sensititre, Trek Diagnostics, Westlake, OH) is used to determine the MICs for 17 antimicrobial agents: amikacin, ampicillin, amoxicillin-clavulanic acid, apramycin, ceftiofur, ceftriaxone, cephalothin, chloramphenicol, ciprofloxacin, florfenicol, gentamicin, kanamycin, nalidixic acid, streptomycin, sulfamethoxazole, tetracycline, and trimethoprim-sulfamethoxazole [Table 2].

Public health laboratories from eight states also select and forward the first *Campylobacter* isolate received each week to CDC for susceptibility testing. For *Campylobacter*, the E-test system (AB BIODISK, Solna, Sweden) is used to determine the MICs for 8 antimicrobial agents: azithromycin, chloramphenicol, ciprofloxacin, clindamycin, erythromycin, gentamicin, nalidixic acid, and tetracycline [Table 2]. After confirmation to genus level, identification of *Campylobacter* to species level is performed using the hippurate test, and, for hippurate-negative *Campylobacter* isolates, and polymerase chain reaction to identify the hippuricase gene, diagnostic of *Campylobacter jejuni*.

For all pathogens in this report, MIC results are dichotomized: isolates with intermediate susceptibility are categorized as sensitive. Breakpoints are determined using, when available, National Committee for Clinical Laboratory Standards (NCCLS) [Table 2].

## Results

### **Non-Typhi *Salmonella***

A total of 1514 non-Typhi *Salmonella* isolates were received at CDC in 1999; 1499/1514 (99%) were viable upon receipt and tested for antimicrobial susceptibility [Table 4a, Figure 1]. Non-Typhi *Salmonella* refers to all *Salmonella* serotypes except serotype Typhi. The antimicrobial agents to which *Salmonella* demonstrated the highest prevalence of resistance were tetracycline, sulfamethoxazole, streptomycin, and ampicillin: 292/1499 (19%) were resistant to tetracycline, 272/1499 (18%) isolates were resistant to sulfamethoxazole, 254/1499 (17%) were resistant to streptomycin, and 234/1499 (16%) were resistant to ampicillin [Figure 2]. Figure 3 provides MIC results for each of the 17 antimicrobials tested. One (0.1%) isolate (*S.* serotype Senftenberg) was resistant to ciprofloxacin; 16 (1%) isolates were resistant to nalidixic acid. Six (0.4%) isolates were resistant to ceftriaxone.



Among non-Typhi *Salmonella* isolates, 388/1499 (26%) were resistant to one or more agents, and 315/1499 (21%) were resistant to two or more agents. Among *Salmonella* isolates tested, 269/1499 (18%) were serotype Enteritidis and 362/1499 (24%) were serotype Typhimurium (includes serotype Typhimurium var. Copenhagen) [Table 5]. In 1999, the serotypes with the highest proportion of isolates which were pansusceptible were Javiana (98%), Thompson (97%), and Braenderup (96%) [Table 6]. Figure 4 provides the resistance among non-Typhi *Salmonella* serotypes from 1996-1999. Among *S. Enteritidis* isolates, 44/269 (16%) were resistant to one or more antimicrobial agents. Among *S. Typhimurium* isolates, 179/362 (49%) were resistant to one or more antimicrobial agents [Table 7]. Figure 5 provides the percent of *S. Typhimurium* by site.

In recent years, a multidrug-resistant strain of *S. Typhimurium* has been identified. This strain is characterized not only by the multidrug-resistant pattern, but also by the phage type – DT104 [Table 8]. Although none of 362 *S. Typhimurium* isolates tested were phage typed, 102 (28%) were resistant to the five antimicrobial agents, ampicillin, chloramphenicol, streptomycin, sulfamethoxazole, and tetracycline (ACSSuT), to which *S. Typhimurium* DT104 is commonly resistant [Figure 6]. Of the 102 *S. Typhimurium* isolates with the ACSSuT resistance pattern, 12 (12%) were also resistant to kanamycin, 9 (9%) were also resistant to cephalothin, 7 (7%) were also resistant to amoxicillin-clavulanic acid, 3 (3%) were also resistant to ceftiofur, and 1 (1%) was also resistant to ceftriaxone [Table 9].

A second penta-resistant pattern, resistance to ampicillin, kanamycin, streptomycin, sulfamethoxazole, and tetracycline (AKSSuT), also has emerged among *Salmonella* Typhimurium [Table 8]. Strains with this resistance pattern are not DT104 by phage typing. Among 362 *Salmonella* Typhimurium isolates tested, 39/362 (11%) had the AKSSuT resistance pattern [Figure 7]. Of the 39 *S. Typhimurium* isolates with the AKSSuT resistance pattern, 12 (31%) were also resistant to chloramphenicol, 8 (20%) were also resistant to cephalothin, and 2 (5%) were also

resistant to amoxicillin-clavulanic acid [Table 9]. Table 10 describes the clinical source of all non-Typhi isolates tested in 1999.

One *Salmonella* isolate (0.1%) was resistant to ciprofloxacin [Figure 3i]. The percentage of *Salmonella* isolates with ciprofloxacin MICs  $\geq 0.25$  increased from 0.4% (5/1326) in 1996 to 1% (15/1499) in 1999 [Table 11]. The percentage of *Salmonella* isolates resistant to nalidixic acid (MIC  $\geq 32$ ) increased from 0.4% (5/1326) in 1996 to 1% (16/1499) in 1999 [Figure 3m]. The percentage of *Salmonella* isolates with decreased susceptibility to ceftriaxone (MIC  $\geq 16$ ) increased from 0.1% (1/1326) in 1996 to 2% (28/1499) in 1999 [Table 12, Figure 3f]. Seventeen of the 97 (18%) *S. Newport* isolates were highly multidrug-resistant, resistant to amoxicillin-clavulanic acid, ampicillin, ceftiofur, cephalothin, chloramphenicol, streptomycin, sulfamethoxazole, and tetracycline as well as having decreased susceptibility to ceftriaxone [Figure 4j].

### **Salmonella Typhi**

A total of 249 *S. Typhi* isolates were received at CDC in 1999; 207/249 (83%) were viable upon receipt and tested for antimicrobial sensitivity [Table 4b, Figure 1]. Of these 207 isolates, forty-one *S. Typhi* isolates were eliminated from analysis because they were duplicate submissions. Among *S. Typhi* isolates, 49/166 (29%) were resistant to one or more antimicrobial agents and 25/166 (15%) were resistant to two or more agents. The most common resistances among *S. Typhi* isolates were to nalidixic acid 31/166 (19%), sulfamethoxazole 28/166 (17%), or streptomycin 23/166 (14%) [Table 3, Figure 8]. Twenty-one (13%) isolates were resistant to ampicillin; 20 (12%) isolates were resistant to chloramphenicol. Figure 9 provides data on *Salmonella Typhi* MICs by antimicrobial agent. None of the *S. Typhi* isolates tested were resistant to amikacin, apramycin, ciprofloxacin, florfenicol, gentamicin, or kanamycin.

## **Shigella**

A total of 377 *Shigella* isolates were received at CDC in 1999; 375/377 (99%) were viable upon receipt and tested for antimicrobial sensitivity [Table 4c, Figure 1]. Among *Shigella* isolates, 341/375 (91%) were resistant to one or more antimicrobial agents and 244/375 (65%) were resistant to two or more agents. The most common resistances among all *Shigella* isolates were to ampicillin 288/375 (77%), tetracycline 215/375 (57%), streptomycin 209/375 (56%), or sulfamethoxazole 206/375 (55%) [Table 3, Figure 10]. One hundred ninety-three (51%) isolates were resistant to trimethoprim-sulfamethoxazole. *Shigella sonnei* isolates were most frequently resistant to ampicillin 219/275 (80%), sulfamethoxazole 150/275 (54%), or streptomycin 143/275 (52%) [Figure 11]. Figure 12 provides data on *Shigella sonnei* MICs by antimicrobial agent. The most common resistances among *Shigella flexneri* isolates were to tetracycline 80/87 (92%), ampicillin 67/87 (77%), or chloramphenicol 56/87 (64%) [Figure 11]. Figure 13 provides data on *Shigella flexneri* MICs by antimicrobial agent. None of the *Shigella* isolates tested were resistant to amikacin, apramycin, ceftiofur, ceftriaxone, ciprofloxacin, or florfenicol.

## **E. coli O157**

A total of 296 *E. coli* O157 isolates were received at CDC in 1999; 292/296 (99%) were viable upon receipt and tested for antimicrobial sensitivity [Table 4d, Figure 1]. Among *E. coli* O157 isolates, 30/292 (10%) were resistant to one or more antimicrobial agents and 12/292 (4%) were resistant to two or more agents. The most common resistances among *E. coli* O157 isolates were to sulfamethoxazole 24/292 (8%), tetracycline 10/292 (3%), and streptomycin 8/292 (3%) [Table 3, Figure 14]. Figure 15 provides data on *E. coli* O157 MICs by antimicrobial agent. None of the *E. coli* O157 isolates tested were resistant to amikacin, apramycin, ceftiofur, ceftriaxone, chloramphenicol, ciprofloxacin, or florfenicol.

## **Campylobacter**

A total of 393 *Campylobacter* isolates were collected in 1999 and forwarded to CDC; 319/398 (80%) were viable upon receipt and tested for antimicrobial susceptibility [Table 4e, Figure 1]. Of the isolates tested, 295/319 (92%) were *C. jejuni*, 20/319 (6%) were *C. coli*, 2 were *C. upsaliensis*, and 2 were *C. fetus* [Table 15].

Among *Campylobacter jejuni* isolates, 158/295 (54%) were resistant to one or more antimicrobial agents, and 60/295 (20%) were resistant to two or more agents. The most common resistances among *Campylobacter jejuni* isolates was to tetracycline 135/295 (46%) followed by nalidixic acid 59/295 (20%), and ciprofloxacin 52/295 (18%) [Table 16, Figure 17a]. Figure 18 provides data on *C. jejuni* MICs by antimicrobial agent.

Among *Campylobacter coli* isolates, 10/20 (50%) were resistant to one or more antimicrobial agents, and 7/20 (35%) were resistant to two or more agents. The most common resistances among *C. coli* isolates was to nalidixic acid 6/20 (30%), tetracycline 6/20 (30%), or ciprofloxacin 6/20 (30%) [Table 16, Figure 17b]. Figure 19 provides data on *C. coli* MICs by antimicrobial agent.

The NARMS 1997-1999 Annual Reports are posted on the NARMS Website. The address is [www.cdc.gov/ncidod/dbmd/narms](http://www.cdc.gov/ncidod/dbmd/narms)

## National Antimicrobial Resistance Monitoring System: Enteric Bacteria 1999 Publications and Presentations

### Publications

1. Zirnstein G, Li Y, Swaminathan B, Angulo F. Ciprofloxacin resistance in *Campylobacter jejuni* isolates: Detection of gyrA resistance mutations by MAMA PCR and DNA sequence analysis. Journal of Clinical Microbiology 1999; 37: 3276-3280.

### Abstracts

1. Dunne E, Fey P, Shillam P, Kludt P, Keene W, Harvey E, Stamey K, Barrett T, Marano N, Angulo F. Emergence of domestically acquired AmpC-mediated ceftriaxone-resistant *Salmonella* serotype Typhimurium (ST) infections. In Program and Abstracts of 39th Interscience Conference on Antimicrobial Agents and Chemotherapy, 1999 September, San Francisco, CA.
2. Fiorentino T, Howard R, Kinney A, Marcus R, Mshar P, Marano N, Westerman J, Reddy S, Angulo F. Routine subtyping of *Salmonella* serotype Typhimurium by PFGE facilitates focused epidemiological investigations in Connecticut. In Program and Abstracts of 39th Interscience Conference on Antimicrobial Agents and Chemotherapy, 1999 September, San Francisco, CA.
3. Fontana J, Bagshaw J, Angulo F, Marano N, Shea D, Goddard A, George H. Plasmid DNA associated with specific bands in PFGE patterns of antibiotic-resistant *Salmonella* serotype Enteritidis. In Program and Abstracts of 39th Interscience Conference on Antimicrobial Agents and Chemotherapy, 1999 September, San Francisco, CA.
4. Hollinger K, Bager F, Marano N, Angulo F, Aaerstrup F, Tollefson L, Gerner-Smidt, Wegener H. Aminoglycoside (AG) resistance in the United States and Denmark: an association between resistance and AG use in food animals, particularly in US poultry sources. In Program and Abstracts of 39th Interscience Conference on Antimicrobial Agents and Chemotherapy, 1999 September, San Francisco, CA.
5. Hollinger K, Silvers L, Marano N, Fedorka-Cray P, Angulo F, Tollefson L, Stamey K. Antibiotic resistance in *Salmonella enterica* Serotypes Heidelberg, Kentucky, and Thompson isolated from human and broiler chicken sources. In Program and Abstracts of 39th Interscience Conference on Antimicrobial Agents and Chemotherapy, 1999 September, San Francisco, CA.
6. Marano N, Benson J, Koehler J, MacKinson C, Wang Y, Madden J, Debess E, Hill B, Archibald L, Boel J, Wegener H, Angulo F. Presence of high-level gentamicin-resistant (HLGR) enterococci in humans and retail chicken products in the US, but not Denmark. In Program and Abstracts of 39th Interscience Conference on Antimicrobial Agents and Chemotherapy, 1999 September, San Francisco, CA.

7. Marano N, Stamey K, Barrett T, Angulo F. High prevalence of gentamicin resistance among selected *Salmonella* serotypes in the US: associated with heavy use of gentamicin in poultry? In Program and Abstracts of Infectious Disease Society of America 37th Annual Meeting, 1999 November, Philadelphia, PA.
8. Marano N, Stamey K, Barrett TJ, Angulo FJ and NARMS: Enteric Bacteria Working Group. The national antimicrobial resistance monitoring system (NARMS): trends in antimicrobial resistance. Emerging Antibiotic Resistance in Foodborne Enteric Pathogens Conference, 1999 August, Athens, GA.
9. Marano N, Stamey K, Barrett TJ, Bopp C, Dabney P, Angulo FJ and the NARMS Working Group. Emerging quinolone-and-extended spectrum cephalosporin-resistant *Salmonella* in the United States.. In Program and Abstracts of American Society for Microbiology, 99th General Meeting, 1999 May, Chicago, IL.
10. Marano N, Stamey K, Barrett TJ, Angulo FJ and NARMS: Enteric Bacteria Working Group. Antibiotic resistance among human *Campylobacter* isolates in the United States, 1997-1998. *Campylobacter, Helicobacter* and Related Organisms Conference, 1999 September, Baltimore, MD.